Automated Commercial Environment—Requirements Recommendation

Date:	July 27, 2001
Number:	ITD-008
Requestor:	ITDS Sub-Committee
Customs Co-Chair:	Don Kusser
Trade Co-Chair:	Tom Anastasi and Sandra Scott

Requirement

FDA: Entry Summary

For Track 4 shipments released without cargo examination, specific commodity information is not submitted until entry summary data is filed. In this case, FDA data for the reported commodities will be collected in the entry summary. The FDA data collected will be similar to that currently collected in ABI, except that ITDS data standards will be adopted to the extent that is possible.

NEXT STEPS: Consult with FDA to incorporate their requirements into the NCAP/P Entry Summary CUSDEC.

Business Need

To maintain integrated reporting of Customs and FDA data, as currently provided by ABI.

Technical Need

A single electronic edit response message reflecting the results of all agencies' edits will be returned. ITDS may need to perform some data conversion in order to support OASIS processing.

Benefits

Permits electronic reporting of data required by FDA. Promotes ITDS goal of fully electronic reporting. Avoids expense of separate FDA reporting.

Risks

Related Subcommittees

Entry

Priority: **Critical** ☐ **High** X **Medium** ☐ **Low** ☐

Customs Use Only

Approved ☐ Not Approved ☐ Further Evaluation Required ☐